

HANULA, P.

Effect of the conditions of wetting on the extract of maize starch. p. 484.

CHEMICKE ZVESTI. (Journal on applied chemistry issued by the Slovak Academy of Sciences and the Slovak Chemical Society. Monthly).  
Bratislava, Czechoslovakia, Vol. 13, No. 7/8, July/Aug., 1959.

Monthly List of European Accessions, (EEAI), LC, Vol. 8, No. 12, Dec. 1959.  
Uncl.

- [illegible]

HANULA, Peter; CUNDERLIKOVÁ, Marta

Production of biologically active yeast by the semicontinuous method. Kvasny prum 10 no. 2: 36-39 F '64.

1. UVUPP, pobočka Bratislava.

SYKORA, M.; HANULIK, J., inz.

Junction rail bonds on the line section at the foot of a hill.  
Zel dop tech 12 no.11:288-289 '64.

HANULOVA, A.

SURNAME (in caps); Given Name

Country: Czechoslovakia

Academic Degrees: /not given/

Institute of Epidemiology and Microbiology (Ustav epidemiologie a mikrobiologie), Bratislava; Director (Riaditel); Doc MUDr J Karolcek

Source: Bratislava, Lekarsky Obzor, Vol X, No 7, 1961, pp 427-437

Data: "Investigations of the Occurrence of Hemolytic Streptococci in Two Children's Collectives in Bratislava."

Authors:

KESTNEROVA, V  
CERVENKA, J

Technical Associates:

HANULOVA, A  
ADAMCOVA, A  
ADAMEC, J

HAN'S, B.

Parallel operation of several regulators, p. 508, STROJIRENSTVI  
(Ministerstvo strojirenstvi) Praha, Vol. 5, No. 7, July 1955

SOURCE: East <sup>E</sup>uropean Accessions List (EEAL) Library of Congress,  
Vol. <sup>4</sup>, No. 12, December 1955

HAUS, E.; TALGA, J.; KIRKLE, M.

An electrical analogy for a heat exchanger.

P. 36. (SLABOPROUDY OZER.) (Praha, Czechoslovakia) Vol. 19, No. 1, Jan. 1958

SO: Monthly Index of East European Accession (EEAI) LC. Vol. 7, No. 5, 1958

HANUS, B.

"ZPA electronic-differential analyzer."

AUTOMATISACE, Praha, Czechoslovakia, Vol. 2, No. 7, July 1959.

Monthly List of East European Accessions (EEAI), LC, Vol. 8, No. 9, September 1959.

Unclassified.



HANUS, Bohumil

General considerations for determining the economy of production lines in making leather ware. Kozarstvi 12 no.12:351-352 D '62.

1. Zavody A.Zapotockeho, n.p., Jaromer.

HANUS, Bohuslav, promovany ekonom, ScC.; DVORAK, Jaroslav, inz.

Some problems of determining the economic effectiveness of continuous transportation. Doprava no.3:223-228 '63.

HANUS, Bohuslav

Simple device for signaling the fallout of phase. Elektro-  
technik 19 no.2:56 F\*64

HANUS, Bohuslav, promovany ekonom, CSc.; DVORAK, Jaroslav, inz.

Problems of determining the economic effectiveness of ropeways.  
Doprava no. 2:89-95 '64.

HANUS, Bohuslav, provovany ekonom CSc.

Normative coefficients of investment effectiveness in  
transportation, and thier determination. Doprava 6  
no.6:408-414 '64.

S/194/62/000/008/028/100  
D201/D308

AUTHOR:

Hanuš, Bořivoj

TITLE:

Problems of optimum control in systems using a  
constant speed servomotor

PERIODICAL: Referativnyy zhurnal. Avtomatika i radioelektronika,  
no. 8, 1962, abstract 82-141 sh (Souhrn prací o auto-  
mat., 1959, Prague, 1961, 77-97 [Czech.; summary in  
Eng.])

TEXT: Analysis of a control system using a servomotor stage with  
either constant or limited speed for any initial conditions. The  
purpose of the analysis is the determination of conditions for mini-  
mum duration of the transient or of programming. The method of suc-  
cessive approximations is proposed for this analysis. The method has  
several advantages over the phase space method. Both methods of so-  
lution are illustrated with concrete examples of automatic control  
systems. [Abstracter's note: Complete translation.]

Card 1/1

HANUS, Borivoj; VAVRA, Karel

Some control algorithms for a digital computer. Stroj na zprac inf  
10:119-139 '64.

1. Zavody prumyslove automatizace National Enterprise, Prague.

HANUS, Borivoj, inz., C.Sc.; STEPAN, Jaromir, inz.

Effect of design parameters on the dynamics of a superheater;  
contribution to the discussion. Automatizace 5 no.7:194 J1  
'62.



HANUS, Borivoj, inz., C.Sc.

Examination of the nonstationary state of a generator. Energetika  
Cz 13 no.4:181-183,190 Ap '63.

1. Zavody prumyslove automatizace, n.p., Praha.

L 12228-63

S/271/63/000/004/042/045

AUTHOR: Hanus, Borivoj

TITLE: A device for converting continuous quantities into discrete quantities, and the reverse

PERIODICAL: Referativnyy zhurnal, Avtomatika, telemekhanika i vychislitel'naya tekhnika, no. 4, 1963, 62, abstract 4B346 (Czechosl. pat., kl. 42m, 14, no. 100824, 15.09.61)

TEXT: The text describes a patented converter for changing continuous quantities into discrete ones, and the reverse; it is characterized by the possibility of using codes of any structure, by the absence of decoding devices and by the property of reversibility. The principle of the invention is based on the fact that the sources of the reference values of the continuous and discrete portions of the device are linked with each other, i.e. mechanically—so that to each value of a continuous reference quantity (voltage or current) there corresponds a definite value of a discrete quantity. The sources of the reference quantities are periodically subjoined to counting devices. One of them is associated with the input of the comparison circuit; to each input of the circuit is delivered a compared signal E. At the moment when the quantities of the compared and the reference signal are equal, from the output of the comparison circuit is delivered a resolving pulse to

Card 1/2

L 12228-63

S/271/63/000/004/042/045

A device for converting .....

the logical circuit; this guarantees receipt of a converted signal at the output of this circuit. For purposes of reverse conversion the sources of reference voltages are changed in position, while the discrete signal subject to transformation is conducted to the comparison scheme. The patented device can be used also as a function generator (continuous or discrete); this makes possible the use of a second source of corresponding signals which realizes the required function or transformation. There are 4 illustrations.

[Abstracter's note: Complete translation]

Card 2/2

HANUS, Borivoj, inz. CSc.

Digital control with limiting conditions on the final control element movement. Automatizace 7 no. 7:169-172 JI '64.

1. Zavody prumyslove automatizace National Enterprise.

L 56451-65

ACCESSION NR: AP5018805

02/0032/64/c114/008/0581/0584

AUTHOR: Hanus, B. (Engineer, Candidate of sciences)

16  
B

TITLE: Approximation modeling of transfer conditions in steam overheaters on analog computers

SOURCE: Strojirenstvi, v. 14, no. 8, 1964, 581-584

TOPIC TAGS: steam superheater, analog computer, computer calculation, approximation calculation

Abstract [Author's English summary, modified]: An explicit calculation of transfer conditions in steam overheaters is extremely difficult, even when an analog computer is used, since the number of available integrators is too low for the system of partial differential equations describing the transfer. The author proposes a method facilitating and shortening calculations by simplifying the mathematical model. Despite its simplicity the method is reliable and was tested in practice.

Card 1/2

L 56451-65

ACCESSION NR: AF5018805

ASSOCIATION: ZPI, Prague

SUBMITTED: 00

ENCL: 00

SUB CODE: IE, DP

NR REF SOV: 000

OTHER: 009

JPRS

Card

282  
2/2

HANUS, Borivoj, inz. BSc.

Control algorithm for digital computers at an arbitrary combination of sampled values of the variables of the system. Automatizace 8 no.1:1-6 Ja '65.

1. Zavody prumyslove automatizace National Enterprise, Prague.

HANUS, Danuta, mgr inz.; KROL, Antoni, mgr inz.

Factors influencing the settlement of slits. Rudy i metale 8 no.3:  
94-97 Mr '63.



HANUS, Danuta, mgr inz.; KRYNICKI, Jerzy, mgr inz.; BONAREX, Edmund,  
mgr inz.

Studies on the nickel sulfate crystallization conditions of  
sulfuric acid solutions. Rudy i metale 9 no. 3:432-437 Ag '64.

HANUS, Danuta, mgr inz.; RIESENKAMPF, Antoni, dr.

Influence of the composition of mixed blends on their hydrometallurgical properties. Rudy i metale 9 nc.12:664-668 D '64.

BONARER Edmund, mgr inz.; HANUS, Danuta, mgr inz.; ZIETV, Jan, mgr

Methods of processing waste electrolytes from copper refineries.  
Rudy i metale 10 no.2:68-72 F '65.

HANUS, F.

5450. UTILIZATION OF LOW GRADE FUELS IN METAL INDUSTRY. HANUS, F. (Hutník (Smelter, Prague), Nov. 1955, vol. 3, 355, 256). An account of recent trends and developments in the utilization of low grade fuel to replace coking coal is given. The use of lignite-charcoal-tar briquettes, the sulphur and ash content of which has been reduced by treatment with hydrogen and chlorine, perfected by Bilkenroth and Rarnier, is described. The need for accurate temperature control when coking low grade coal, and the disadvantages of quenching coke with water, the use of charcoal and anthracite-coke mixtures in Russian blast furnaces, and developments in fuel saving techniques in Czechoslovakia are considered.

I.S.I.

HANUS, J.

"First automatic fish lift at the Nosice Water Works." p. 154.

STAVBA. (Poverenictvo stavebnictva). Bratislava, Czechoslovakia,  
Vol. 6, No. 5, May 1959.

Monthly list of East European Accessions (EEAI), LC, Vol. 8, No. 8,  
August 1959.  
Uncla.

HANUS, J.

Function and organization of designing in an enterprise. p. 628

POZEMNI STAVBY. (Ministerstvo stavebnictví) Praha, Czechoslovakia, Vol. 7, no. 12, 1959

Monthly List of East European Accessions (EEAI), LC. Vol. 9, no. 2, Feb. 1960

Uncl.

HANUS, J. - Strojirenetvi - Vol. 5, no. 4, Apr. 1955.

Mathematical machines help industry. p. 306.

SO: Monthly list of East European Accessions. (REAL), LC, Vol. 4, No. 9, Sept. 1955  
Uncl.

HANUS, J.

HANUS, J. Sidecar for the JAWA-CZ motorcycle. p. 368

Vol. 10, no. 12, June 1956  
SVET MOTORU  
TECHNOLOGY  
Praha, Czechoslovakia

So: East European Accession Vol. 6, no. 2, 1957



HANUS, J. Ing. (Praha XII, Tr. W. Piecka 98)

New method of planning preparation. Pracovní lek. 9 no.4:350-352  
Sept 57.

1. Ministerstvo zdravotnictví, hygienický a protiepidemický odbor.  
(PUBLIC HEALTH,  
in Czech., planning of hyg. & anti-epidemic center (Cz))

HANUS, J.

Jaroslav Hanus and Vladimir Munk, "Metabolismus von Kohlenhydraten bei Eremothecium Ashbyii," Die Naturwissenschaften (Berlin), 45/1, January 1958, p. 14.

Received on 14 November 1957.

Research Institute for Food Technology, Prague 16, Na Belidle 21.

HANUS, J.; MUNK, V.

7th International Microbiological Congress in Stockholm. p. 298.

PRUMYSL POTRAVIN. (Ministerstvo potravinarskeho prumyslu) Praha, Czechoslovakia,  
Vol. 10, no. 6, June 1959.

Monthly list of East European Accessions (EEAI) LC, Vol. 8, No. 11,  
November 1959.

uncl.

Statistical control in the food industry. p. 506

Statistical control in the food industry. p. 506

PRŮMYSL POTRAVIN. (Ministerstvo potravinářského průmyslu)  
Praha, Czechoslovakia Vol. 10, no. 10, Oct. 1959

Monthly List of East European accession, (EAT), 10, Vol. 2, No. 12, Dec. 1959  
Uncl.

CZECHOSLOVAKIA/Chemical Technology. Chemical  
Products and Their Applications.  
Food Industry.

H

Abs Jour : Ref Zhur-Khimiya, No 6, 1959, 21285

Author : Hanus, Jaroslav; Matejovsky, Karel;  
Necadova, Marie

Inst : -

Title : Sanitation in Bread Baking.

Orig Pub : Prumysl potravin, 1958, 9, No 6, 317-321

Abstract : No abstract.

Card : 1/1

H-123

HANUS, Josef

DECEASED

*Chemistry*

see ILC

HANUS

Y. HANUS

and the calculation of the capacitance of the bushings

SA

5909. Graphical calculation of condenser bushings considering a finite number of metal foils. (HANUS, Y. *Elektrotech. Obz.*, 37, 465-70 (Nov., 1948) In Czech.—Earlier work by the author (*Elektrotech. Obz.*, 165 (1943)) gave a nomographic method for determining the relation between ratio of the reduced thicknesses of the first and last insulating layers, and the ratio of lengths of the first and last metal foils. Uniform voltage distribution in an axial direction and a finite number of foils were assumed. This solution considers the expression for the max. radial gradient between two consecutive conducting cylinders, i.e. the gradient on the surface of the inner cylinder,  $G_r = 2E/(D_{K-1} \log D_K/D_{K-1})$ . This is more exact than the formula for an infinite number of foils, the accuracy of which is not sufficient for  $< 10$  foils. The present paper enables the method to be applied in practice, giving the min. dia. of bushing for a given conductor diameter, the length ratios of the conducting cylinders, the coefficients entering the formula, etc.

B. F. K.

B 64

Hanus, J.

621.314.224  
✓ 5051. Short-circuit-proof current transformers. J.  
HANUS. *Elektrotech. Obozr.* 44, No. 3, 286-90 (1955)  
EE In Czech.

The capacity of current transformers to withstand short-circuits is an important characteristic in the case of branch lines of comparatively small power input supplied by installations of high output. A nomogram is given for selecting the degree of short-circuit-proofness of a current transformer to suit given conditions. Two production series of such transformers, for cascade connection, with overcurrent factors of 250-1 300 are described.

ELECTRICAL RESEARCH ASSOCIATION

Handwritten initials: "JH" and "MT" with a checkmark.



HANU, J.; KOTICEK, J.

Current and voltage transformers for 220 kV. p. 306.

ELEKTROTECHNIK. Vol. 11, no. 10, Oct 1956

Praha, Czechoslovakia

SOURCE: East European List (EEAL) Library of  
Congress, Vol. 6, No. 1, January 1957

HANUS, J.

2

631.314.232  
2124. INCREASE OF THE OUTPUT OF VOLTAGE TRANS-  
FORMERS BY AN AUXILIARY COMPENSATING DEVICE.

J. Hanus, "Elektrotech. Obzor," Vol. 45, No. 2, 57-63 (1956). In Czech. (1)

The principle of the compensating circuit for a voltage transformer is explained and the practical application described with reference to the type NAK/COF 110 compensator. An elementary theory of the errors of voltage transformers is given, together with a method of determining the components of the transformer impedance from results of the calibration. The increase in output obtained range from 130 to 280%.

Electrical Research Association

*[Handwritten signature]*

HANUS, J.

TECHNOLOGY

ELEKTROTECHNICKY OBZOR.

HANUS, J. Answering the discussion concerning the terms amperzavity and proudeni  
p. 657.

Vol. 47, No. 12, Dec., 1958

Monthly List of East European Accessions (EEAI) LC, Vol. 8, No. 5 May 1959, Unclass.

S/196/62/000/022/007/007  
E194/E155

9.2/20  
AUTHORS: Hanus, Jan, and Vrtel, Leo

TITLE: Cascade current transformer

PERIODICAL: Referativnyy zhurnal, Elektrotehnika i energetika,  
no.22, 1962, 28-29, abstract 22 I 149 P. (Czech. pat.  
cl. 2ld<sup>2</sup>, 53/01, no.99246, April 15, 1961)

TEXT: The bushing-type current transformer, the construction of which is patented, withstands short-circuit currents very well. It is a two-stage cascade. The construction is similar to that of a pedestal-type current transformer, which is simpler, because in it the short-circuit current passes only through the primary winding of the first stage, whereas in that under consideration the current must also pass through the second stage. Only the second stage is insulated for high voltage. The first stage has low-voltage insulation, so that its core (made of toroidal strip) has a very short mean magnetic path length and consequently low magnetising ampere turns. For a given accuracy this makes it possible to use a small rated number of ampere-turns in the first stage. The primary winding (2) and the secondary winding (3) of  
Card 1/5

Cascade current transformer.

S/196/62/000/022/007/007  
E194/E155

the first stage (see sketch) are uniformly distributed over the first core. The small number of uniformly-distributed circular turns and the small dimensions of the core give the first stage rigidity against short-circuit currents. The second stage has two primary windings (5), (6) and secondary winding (7). Its core (4) may be toroidal, or even rectangular of core or shell-type. The primary winding (5) is a continuation of winding (2) of the first stage, and short-circuit current also passes through it. For rigidity, it should have the least possible number of turns, preferably one. The conductor section of windings (2) and (5) should be designed to ensure thermal stability. A further second stage primary winding (6) is connected to the second winding of the first stage (3). The considerable burden which the second stage represents, and also the small section of the primary stage core, causes this core to saturate when the current is heavy. Therefore, the current in windings (3), (6) and (7) is considerably less than in windings (2) and (5) (it does not exceed 50 times the rated current). The section of conductors for windings (3), (6) and (7) should be selected accordingly. Windings (5) and (6) should be

Card 2/5

Cascade current transformer

S/196/62/000/022/007/007  
E194/E155

wound in such a way that within the core currents in them are in the same direction. The number of turns in winding (7) is based on the condition

$$I_2 N_2 = I_1 N_1 + I_1^1 N_1^1, \quad \text{where } I_1, I_1^1 \text{ and } I_2 \text{ are}$$

the currents in windings 5, 6 and 7, and  $N_1, N_1^1$  and  $N_2$  are the number of turns in these windings. Since windings 5, 6 and 7 need only normal rigidity against short-circuit currents, the second stage may consist of ordinary bushing-type current transformers with porcelain insulators (with somewhat modified winding data). To these current transformers is fitted a first stage the data of which depend on the rated current, the rigidity class and the short-circuit current. Possible variants are: 1) winding (5) may have one or several turns; 2) the first stage may be made as an auto-transformer with a transformation ratio of 1:1; 3) to reduce the error, the first stage may be made with pre-magnetisation - the first stage core is divided into two unequal parts, the smaller having a premagnetising winding supplied from the terminals of an impedance connected in series with winding (6);

Card 3/5

Cascade current transformer

S/196/62/000/022/007/007  
E194/E155

4) if the second stage core is of the shell type it is not uniformly magnetised because winding (5) passes through only one of the two windows of the core; to avoid this an equalising winding is wound on the adjacent part of the core and connected in series with winding (7); the number of turns in this equalising winding,  $N_2^1$ , is selected according to the condition  $I_1 = I_2 N_2^1$  (supposing that winding (5) has one turn). The advantages of the construction are: high rigidity; no need for ferro-nickel alloys as in single-stage construction; the possibility of using standard multi-turn bushing transformers for the main part of the current transformer.

[Abstractor's note: Complete translation.]

Card 4/5

38036

Z/017/62/051/006/002/003  
D409/D301

9.2120

AUTHOR: Hanuš, Jan, Engineer, Doctor

TITLE: Evaluating the economy of error-compensation in an instrument current transformer

PERIODICAL: Elektrotechnický obzor, v. 51, no. 6, 1962, 266-276

TEXT: The article describes three cases of transformer coupling used for error compensation in instrument transformers produced by the Elektrotechnické Závody J. Fučíka (J. Fučík Electro-technical Engineering Plant) in Brno. In all three cases, the magnetic circuits of the current transformer were excited from an external source with a current of three times the nominal frequency. In one of the cases, this pre-excitation was combined with the compensation of electromotive force in the turns of the main magnetic circuit and the compensation of the no-load component of the excitation current with the aid of a capacitor. The effect of this compensation is evaluated by derivation of the specific admittance, dependent on the specific electromotive force, from the errors of a non-

Card 1/3



Evaluating the economy ...

Z/017/62/051/006/002/003  
D409/D301

compensated and a compensated transformer. Admittance characteristics are then compared with each other and with the characteristics of commonly used ferromagnetic materials, giving a comprehensive survey on the economy of the compensation method used. This economy may be judged according to how much and what type of ferromagnetic material, coupled with the primary winding, can be saved by the compensation. At given values of the mean line-of-force length of the nominal flow, and the demands imposed upon output and accuracy, the cross-section area of the magnetic circuit is directly proportional to the specific admittance of the ferromagnetic material used. This specific admittance can be reduced on instrument current transformers by various methods of compensation. Results obtained by pre-excitation, as described in the article, and relating to hot-rolled, or cold-rolled, magnetically oriented silicon steel transformer sheets (sonaperm), show that under identical conditions it is possible to reduce by one half the cross-section area of the magnetic circuit as compared with transformers not pre-excited. The applied notion of specific admittance and specific voltage for constant frequency is more advantageous when calculating current-

Card 2/3

Evaluating the economy ...

Z/017/62/051/006/002/003  
D409/D301

transformer errors than the notion of permeability. In conjunction with values for reduced impedances, load and loss, and further flow consideration, one thus obtains a more distinct outline of the laws of transformer growth than by considerations based on magnetic inductance and permeability of the ferromagnetic material. There are 10 figures and 5 tables. The English-language references are: A. Boyajian: Orthomagnetic Bushing Current Transformer for Metering. Transaction AIEE (1945), pp. 137-140; A.C. Schwager: Current Transformer Performance Based on Admittance Vector Locus. Transaction AIEE (1942), pp. 26-30. (Technical editor: J. Kopeček). 4

ASSOCIATION: ŘOS-EJF Brno

SUBMITTED: March 10, 1962

Card 3/3

MUNK, V; PASKOVA, Jirina; HANUS, J.

Factors Influencing Glucose Oxidase Activity in Submerged Cultivation  
of *Aspergillus niger* on Synthetic Medium. Folia microbiol. 8 no. 4:203-14  
Jl '63

Department of Microbiology, Central Research Institute of the Food  
Industry, Prague.

(OXIDOREDUCTASES) (ASPERGILLUS) (CULTURE MEDIA) (CITRATES)  
(GLUCOSE) (CALCIUM) (MAGNESIUM) (AMMONIA)  
(HYDROGEN-ION CONCENTRATION)

HANUS, Jaroslav, inz. RNDr., CSc.; HOJDEM, Bretislav, doc., inz.

Enzymatic determination of glucose in biological materials.  
Prum potravin 14 no.10:552-554 0 '63.

1. Ustredni vyzkumny ustav potravinarskeho prumyslu, Praha (for Hanus). 2. Universita 17.listopadu, Praha (for Hojdem).

HANUS, K.

Two metabolic levels in bats. Cesk. fysiол. 7 no.3:232-234 May 58.

1. Biologická fakulta KU, Praha.

(METABOLISM,  
curves in bats (Cz))  
(BATS,  
metab. curves (Cz))

CZECHOSLOVAKIA

AUS, ...; Laboratory of Physiology and Genetics of Animals,  
Czechoslovak Academy of Sciences (Laborator Fysiologie a Genetiky  
Zivocichu CSAV), Uhřetín.

"Investigation of Thermoregulation in Ruminants."

Prague, Ceskoslovenska Fysiologie, Vol 15, No 5, Sep 66, pp  
374 - 375

Abstract: The determination of basal metabolism in ruminants is discussed. During the first few days of starvation the metabolism at rest does not decrease, and shows a slightly increasing tendency, reaching a maximum on the 3rd day. The correct way of measuring the metabolism of ruminants is within the first 24 hours of starvation. In sheep the measurement should be taken with the wool sheared to 3 mm; because of the excellent insulating properties of wool misleading results would be obtained otherwise. 1 Figure, no references. Submitted at 3 Days of Physiology of Domestic Animals at Liblice, 8 Dec 65.

1/1

"APPROVED FOR RELEASE: 09/19/2001 and CIA-RDP86-00513R000617910003-4"  
Czechoslovak Academy of Sciences (Laborator Fysiologie a Genetiky  
Zivocichu CSAV), Uhřetín.

"The Technique of Determination of Total Metabolism of Sheep."

Prague, Ceskoslovenska Fysiologie, Vol 15, No 5, Sep 66, pp  
375 - 376

Abstract: The author describes an inexpensive installation which he designed for the measurement of metabolism in sheep. A diagram and description of the installation are presented. Methods of heat removal and air supply are discussed. Thermostatic instrumentation can be regulated to maintain temperatures of  $-10^{\circ}$  to  $+40^{\circ}\text{C}$ . The accuracy of the thermostat is  $\pm 1^{\circ}\text{C}$ . Overall dimensions of the installation are 4x3x2 meters. 1 Figure, no references. Submitted at 3 Days of Physiology of Domestic Animals at Liblice, 8 Dec 65.

1/1

HANUS, L.

Terraces of the Mze River between Tachov and Krimice in the vicinity of plzen. p.81.  
(Casopis Pro Mineralogii A Geologii, Vol. 62, No. 2, 1957, Praha, Czechoslovakia)

SO: Monthly List of East European Accessions (EEAL) LC. Vol. 6, No. 9, Sept. 1957. Uncl.

HANUS, M.

Optimum reaction conditions for preparation of 2-vinylfuran. I. Šimek and M. Hanuš (Slovenská vysoká škola tech., Bratislava, Czech.) ~~Chem. zvesti 13, 108-10 (1959)~~ (German summary).—The effect of reaction time, temp. of the bath, amt. of  $\text{Ac}_2\text{O}$ , and of anhyd.  $\text{AcONa}$  on 1 mole furfural and catalysts  $\alpha$ -picoline, pyridine, dimethylaniline, and  $\text{K}_2\text{CO}_3$  on the Perkin synthesis of  $\beta$ -(2-furyl)acrylic (I) acid was studied. It was detd. that on decarboxylation of I without quinoline, the effect of the decomp. acid and the catalyst was in the area of exptl. error. By this modified method it was possible to increase the yield of raw 2-vinylfuran by 10-15% as compared with the data in the literature.

Jan Micks

2 May  
4E 2c gj  
4E 3d  
4

gJ



HANUS, M.

Distr: 4E2c(j)

/ Polymerization of 2-vinylfuran. I. Simek and M. Hanus (Slovenská vysoká škola tech., Bratislava, Czech.). Chem. zvesti 14, 124-8 (1960) (German summary).—By block polymerization of 2-vinylfuran in N atm., elastic polymers of light color, sol. in org. solvents can be obtained. The polymerization speed is relatively small. The application of strong inorg. acids and  $SiCl_4$  and a thermal polymerization at higher temps. ( $150^\circ$ ) form hard, shiny, dark colored, insol., and infusible polymers. The speed of emulsion polymerization initiated by Na persulfate was detd. at  $17-60^\circ$ . Emulsion polymerization can be carried out at temps. considerably higher than reported in literature, and by increasing temp. up to  $50^\circ$ , the time of the polymerization can be shortened to approx.  $\frac{1}{10}$  (from 48 to 5 hrs.). Jan Micka—

4  
KJMS (HUB)  
1-3 HI (DRY)

HANUS, Milan; VACLAVIK, Frantisek

Ensuring the winter operation of the Czechoslovak Airlines.  
Letecky obzor 6 no.11:350-351 '62.

HANUS, Rudolf, inz.

Tests of a dual teletype line. Cs spoje 7 no.11:13-14 N '62.

1. Jihomoravská krajská správa spoju.

CZECHOSLOVAKIA/Chemical Technology. Chemical Products and  
Their Applications. Chemical Engineering.

H

Abs Jour: Ref Zhur-Khim., No 8, 1959, 27696.

Author : Manus, S.; Kolar, V.

Inst :

Title : Study of Mixing. II. Analytical Method Using Insoluble  
Particles; Statistical Experimental Method for the Study  
of Mixing in Liquid Media. III. Effect of Mechanical  
Agitation on Rate of Solution of a Granular Solid Phase.

Orig Pub: Chem Listy, 52, No 5, 839-851, 852-858 (1958) (in Czech)

Abstract: II. An experimental method has been developed for the  
study of a mixing process based on the observation of  
a single particle in the agitated liquid. The method  
permits the investigation of mixing conditions in var-

Card : 1/3

130

CZECHOSLOVAKIA/Chemical Technology. Chemical Products and  
Their Applications. Chemical Engineering.

H

Abs Jour: Ref Zhur-Khin., No 8, 1959, 27696.

ious portions of the liquid and thus makes it possible to obtain a more detailed picture of the mixing taking place in the space occupied by the agitated liquid. The data obtained by this method are in agreement with data obtained by other workers.

III. An equation has been derived for the rate of solution of a granular solid phase in a mechanically agitated liquid. The equation expresses the Nusselt diffusion number as a function of the Reynold's number, the Schmidt number, and two simplex parameters, one of which reflects the effect of the specific gravity of the two phases, and the other gives the effect of tur-

Card : 2/3

CZECHOSLOVAKIA/Chemical Technology. Chemical Products and  
APPROVED FOR RELEASE: 09/19/2001 CIA-RDP86-00513R000617910003-4  
Their Applications. Chemical Engineering.

Abs Jour: Ref Zhur-Khin., No 8, 1959, 27696.

bulence. For Communication I see RZhKhim, 1958, 60803. -- M. Ryba.

Card : 3/3

RIBAL, Miloslav, inz.; HANUS, Stanislav, inz.; BERNASEK, Vladimír

Use repellents as protection against damage caused by deer. Agrochem  
2 no.1:18-22 '62.

1. Československá akademie zemědělských věd, Vězkumny.ustav lesního  
hospodářství a myslivosti, Zbraslav (for Ribál and Hanus). 2. Spolana,  
Neratovice (for Bernasek).

DOLEJS, L.; HANUS, V.

"Mass spectrometry; organic chemical applications" by  
K. Biemann. Reviewed by L. Dolejs, V. Hanus. Coll Cz  
Chem 29 no.4:1083 Ap '64.

HANUS, VACLAV

✓ *Ger* Mineralogy and geochemistry of the copper-lead-zinc deposit of Vrančice, with chalcocite and willemite (Příbram ore region, Bohemia). Václav Hanus. *Sborník ústřed. ústavu geol.* 22, 69-144 (1955) (English summary).—The presence of chalcocite and willemite is unusual in primary ore deposits. A study of succession shows that they were formed in a process of hypogene mineralization. The main minerals found are sphalerite, galena, and hematite. The gang consists mainly of calcite and dolomite. The ore deposit shows signs of three periods of deposition. The ores were deposited in the following order: Zn-Pb-Cu, Ag-As, Sb. The deposit is characterized by a S deficiency, as indicated by the presence of native Cu and Ag and of Fe oxides.  
R. Pick

L



HANUS, V.

Problems of the deposit-forming metasomatism. In German. p. 74

Prague. Ustredni ustav geologicky. VESTNIK. Prague, Czechoslovakia, Vol. 34, no. 1, 1959

Monthly List of East European Accessions (EEAI), LC, Vol. 8, no. 11, Nov. 1959  
Uncl.

BERNARD, Jan H.  
SURNAME, NAME

Country: Czechoslovakia

Academic Degrees: /not given/

Affiliation: Central Geological Institute (Ustredni ustav geologicky), Prague.

Source: Prague, Vestnik Ustredniho Ustavu Geologickeho, Vol XXVI, No 5, June 1961, pp 343-345. 361-363.

Data: "The Position of Hematite (Specularite) and Magnetite in the General Succession of the Hydrothermal Deposits of the Spissko-Gemerske Rudohorie Mountains."

"The Time Relation of the Gemerid Granites and Tourmalization to the Hydrothermal Mineralization in the Spissko-Gemerske Rudohorie Mountains."

Authors: BERNARD, Jan H.

HAJOS, Vaclav

HANUS, V.



CZECHOSLOVAKIA

HANUS, V; KRS, M.

1. Central Geological Institute (Ustredni ustav geologicky),  
Prague ; 2. Institute of Geophysics (Ustav uzite  
geofyziky), Prague (for all)

Prague, Vestnik ustredniho ustavu geologickeho, No 2, 1963,  
pp 119-121

"Paleomagnetic Verification of the Neoidic Age of Hydro-  
thermal Mineralization in the Krusne Hory Mts.  
and in the Slavkovsky les."

HANUS, Vaclav, promovany ekonom.

Labor economy and labor biological problems. Prace mzda 11  
no.4:189-192 Ap '63,

HANUS, Vaclav, promovany ekonom

Work remuneration and biological labor problems. Prace mzda 11  
no.5:235-238 My '63.

HANUS, Vaclav ; KRS, Miroslav

Palaeomagnetic dating of hydrothermal mineralization on example of Spissko-Gemerske rudohorie area, Czechoslovakia. Rozpravy met CSAV 73 no.14:1-87 '63.

1. Central Geological Survey, Prague 1, Hradelni 9; Institute of Applied Geophysics, Prague 5, Podbelohorska 47.

41-3485, ✓  
24(2,4)

PHASE I BOOK EXPLOITATION

CZECH/2433

International Polarographic Congress. 1st, Prague, 1951

Sporník I. Mezinárodního polarografického sjezdu. Díl 3: Hlavní referáty přednesené na sjezdu. Proceedings...Vol 3: Reviews Read at the Congress. Praha, Přírodovědecké vyd-ví [1952] 774 p. 2,000 copies printed.

Resp. Ed.: Jiří Koryta, Doctor; Chief Ed. of Publishing House: Milan Skalník, Doctor; Tech. Ed.: Oldřich Dunka.

PURPOSE: The book is intended for chemists, chemical engineers, and physicists.

COVERAGE: The book is a collection of reviews and original papers read at the International Polarographic Congress held in Prague in 1951. Uses of polarography in organic and inorganic analysis, biochemistry, medicine, and industrial chemistry are discussed. In this section, Reviews Read at the Congress, Russian and either German or English translations of each review are presented. In the section, Original Papers Read at the Congress, only those translations in Russian, German, and English which

Card 1/14

Proceedings (Cont.)

CZECH/2433

have not been published in Volume I are presented. The following scientists participated in the opening of the Congress: Professor Wiltor Kemula, Dean of the Faculty of Sciences, Warsaw; Doctor Jaromir Dolansky, Minister of Planning; Professor Jaroslav Herovsky, Chairmen of the Congress; and Professor Jaroslav Fukatko, Chairman of the Center for Scientific Research and Technical Development. References follow each paper.

TABLE OF CONTENTS:

REVIEWS READ AT THE CONGRESS

Heyrovsky, J. Fundamentals of Polarography	5
[Russian Translation]	13
[English Translation]	22
Majer, Vl. Polarography in Inorganic Analysis	32
[Russian Translation]	55
[German Translation]	80
Hanus, Vl. Polarographic Behavior of Organic Compounds	103
Card 2/14	



Proceedings (Cont.)	CZECH/2433
[Russian Translation]	118
[English Translation]	132
Zuman, P. Organic Analysis	145
[Russian Translation]	160
[German Translation]	177
✓ Santavy, F. Polarography in Biochemistry and Medicine	194
[Russian Translation]	210
[German Translation]	226
Forejt, J. Apparatus for Oscillographic Polarography	241
[Russian Translation]	250
[German Translation]	259
Heyrovsky, J. Oscillographic Polarography	268
[Russian Translation]	273
[English Translation]	279
✓ Brdicka, R. Kinetics of Electrode Processes in Polarography	286
Card 3/14	

Proceedings (Cont.)

CZECH/2433

Svatek, E. Study of Catalytic Reactions at a Dropping  
Mercury Electrode

667

Koryta, J. Decomposition Rate of the Complex of  
Nitrilotriacetic Acid With Cadmium

672

Smutek, M. Slow Electrode Reactions

677

[Russian Translation]

683

[English Translation]

687

Hanus, V. Polarographic Study of the Recombination of  
Phenylglyoxylic Acid

691

Koutecky, J. Linear Systems of Electrode Reactions in  
Which a Chemical Reaction in Solution Takes Place

699

Pliva, J. Contribution to the Theory of Diffusion Cur-  
rents

708

[Russian Translation]

712

[English Translation]

717

Card 12/14

CA  
HABUSH, [V.]

Polarographic study of the decomposition of penicillin in acidic solutions. V. Habuš and E. Křečí (Charles Univ., Prague, Czech.). *Chem. Listy* 46, 52-3(1952).— Benzylpenicillin is reduced polarographically at pH 4.6

giving 2 waves. The max. diffusion current is found at pH 2.2. The wave height decreases with increasing time of reaction. Benzylpenicillic acid is believed to be the proper reducing component. The polarographic detn. of penicillin is suggested. M. Hudlický

CA  
HANUSH, V.

The effect of the ionic recombination on the polarographic reduction of dibasic acids. A. Hama and R. Hudlický (Charles Univ., Prague). *Chem. Listy* 44, 291 (1950). Equations have been derived which account for the effect of ionic recombination on the polarographic limiting currents caused by the reduction of the undissociated molecules belonging to dibasic acids. As long as the limiting reduction currents of the undissociated molecules appear in the range of pH where these molecules are practically absent in the body of the solution, their variation with pH can be expressed as follows:  $2 \text{ pH} = \text{pK}_1 + \log \frac{i_{\infty} - i}{i_{\infty} - i_{\infty}} + \log \frac{i_{\infty}}{i_{\infty}}$ . Here,  $i_{\infty}$  denotes the limiting reduction current of univalent anions,  $K_1$  the 2nd acid dissociation constant, which is to the first apparent polarographic dissociation constant of the specific rate because of the ionic recombination and of the drop time owing to the capillary electrode. The equation concerned shows that the apparent polarographic dissociation curve for dibasic acids is not, in general, a symmetric dome, and it can be twice as steep as the dissociation curve owing to univalent acids. All the relations derived theoretically have been verified by experimental data obtained with malonic, oxalonic, and fumaric acids. The thickness of the reaction layer surrounding the surface of the capillary electrode (according to the Wiener statistical conception) and the absolute rate constant of ionic recombination for the above acids have been computed.

M. A. Hudlický

1951

CH HANUSH, [V.]

Electrochemistry - 9

Polarographic study of the recombination of phenylglyoxylic acid. VI. HANUSH (Charles Univ., Prague). *Sbornik Mezinárodn. Polarog. Sjezdu Praha, 1st. Congr. 1951, Pt. 1, Proc. 801-11 (in Russian), 811-17 (in German).* — The reduction of phenylglyoxylic acid is irreversible over the pH range 0-14. In very acid and very alk. solns. only one wave is obtained which is ascribed to the undissoc. mole. and to the corresponding anions, resp. The 2 types of reductions occur simultaneously over the range of pH 5-8, resulting in 2 waves. The first of these is largely detd. by the velocity of the recombination of anions with  $H^+$  at the surface of the dropping-Hg electrode. The ratio of this kinetic current (1st wave) to the total current (sum of both waves) was studied in detail. It was independent of the concn. of the depolarizers and was not influenced by deformable cations or anions, buffer acid, and capillary active materials, nor by a change in the resistance of the elec. circuit. It is only little influenced by the velocity of outflow of Hg and by temp. However it is mainly controlled by pH and drop time as well as by the concn. of the buffers and the ionic strength of the soln. Certain deviations between the exptl. data and calcn. based on the equations of Koutecký and Brlička (*C.A.* 43, 484) are discussed. O. H. Müller

Hanus, V.

Evaluation of the rate constant of the decomposition of hydrogen peroxide by catalase from the polarographic limiting current of oxygen. J. Koutecký, R. Hudský, and V. Hanus (Czech. Acad. Sci., Prague). Collection Czech. Chem. Commun. 18, 611-28(1953)(in English), Chem. Listy 47, 763-805(1953).--Previous interpretations of the effect of catalase on the polarographic limiting currents due to the reduction of O were revised. Calculations of the rate constant, based on the concept of a reaction layer at the electrode were of only limited value. A rigorous solution was given of the appropriate system of differential equations, taking into account the growth of the Hg drop, for a reaction in which half the originally present reactant (O) was regenerated from the reduction product ( $H_2O_2$ ) by a chem. reaction with the catalyst (catalase). Values for the resulting function were tabulated, and the rate constants, calculated by means of it, were in good agreement with values obtained by direct manometric measurements.

Otto H. Müller

<sup>H</sup>  
Hanus, V.; Koutecky, J.; Briedka, R.

Hanus, V.; Koutecky, J.; Briedka, R. "Calculation of the rate constant for the decomposition of hydrogen peroxide by catalase from polarographic limiting currents caused by oxygen. p. 793. CASOPIS PRO FESTOVANI MATEMATIKY. CZECHOSLOVAK MATHEMATICAL JOURNAL. Vol. 47 No. 6 June 1953 Praha, Czechoslovakia.

SO: Monthly List of East European Accessions, L. C. Vol. 3 No. 1 Jan. '54 Uncl.

*Hanus, V.*  
Czechoslovakia/Physical Chemistry. Electrochemistry

B-12

Abs Jour : Ref Zhur - Khimiya, No 7, 1957, 22519.

Author : Vladimir Hanus.

Inst : Not given

Title : Solution of Problems Related to Polarographic Kinetic Points, Limited by Bi-molecular Chemical Reactions with the Aid of Approximation Method.

Orig Pub : Chem. zvesti, 1954, 8, No 10, 703-713. (czech., Res. russ., nem.)

Abstract : Bases of approximation method (of Brdichki and Visner) of solutions of problems of kinetic current limitations in polarography are stated, in which it is assumed that a) chemical reaction takes place in a thin near-electrode layer of a determined extent; b) substance concentrations, taking part in the process, have certain average values which do not change during the life of a drop; c) concentration gradient of the electroactive form on the surface of an electrode is equal to the difference of concentrations on the surface and at the limit of the reactive layer divided by the thickness of the reactive layer; d) concentration of inactive forms at the surface of the electrode is equal to the difference between the hypothetically dif-

Card 1/2

-183-

Czechoslovakia/Physical Chemistry. Electrochemistry.

**APPROVED FOR RELEASE: 09/19/2001** **CIA-RDP86-00513R000617910003-4**  
Abs Jour : Ref Zhur - Khimiya, No 7, 1957, 22519.

fused current which would occur at an infinite high speed of the chemical stage, and the observed kinetic current divided by the constant of Il'covich's equation for the given form. Six examples of equations for kinetic currents are evolved in the course of 2nd degree reactions on the electrode. It is indicated that the method of approximation permits to resolve easily problems insoluble or difficultly soluble by a precise method, but speed constant values thus obtained could be 2 or 3 times smaller or larger than the actual ones.

Card 2/2

-184-

*HANUS, VLADIMIR*



HANKS, V

CZECH

621.316.722.9

3146. A degeneration voltage stabilizer with a possibility of overcompensation. M. PACAK AND V. HANUS. *Slahoproudy Obzor*, 19, No. 2, 75-82 (1955) in Czech.

The system is particularly suitable for stabilization of voltages of the order of several kV, since its d.c. amplifier consists of two valves operating with a common cathode resistance. A source of reference voltage can be connected into the grid circuit of either of the amplifying valves. Operation of the circuit and its stability are analysed in detail. Two practical stabilizers (2.0 kV and 200 V) are described in detail, their performance being illustrated by a number of experimental curves. The system is compared with a stabilizer designed by R. B. Mackenzie [Abstr. 976 (1954)].

R. S. Sidorowicz

1

11/10/55

HANUS, VLADIMIR

CZECH

Two examples of rapid bimolecular reactions participating in the depolarization process on the dropping mercury electrode. Jiroslav Koucky and Vladimir Hanus. *Collection Czechoslov. Chem. Commun.*, 20, 134-38 (1955) (in German). See C.A. 49, 14451. E.J.C.

14X 224

~~Vermine~~ Hanus, VE.

CZECHOSLOVAKIA/Nuclear Physics

C-2

Abs Jour : Referat Zhur - Fizika, No 5, 1957, 11001

Author : Cermah Vladimir, Hanus Vladimir

Inst : Not given

Title : Remarks on the Article by Ondracek "Mass Spectrograph of the Research Institute of Vacuum Electronics."

Orig Pub : Slaboproudy obzor, 1956, 17, No 8, 456-457

Abstract : See Referat Zhur Fizika, 1956, 1583.

Card 1/1

HH105, V.  
Polarographic behavior of phthalic acid and some of its  
esters. A. Ryvolová and V. Hanuš, *Collection Czech.  
Chem. Commun.* 21, 853-61 (1956) (in German).—See C.A.  
50, 4675f.

2  
PM

HANUS, VLADIMIR

5

✓ Polarographic behavior of phthalic acid and some of its esters. Anna Ryvolová and Vladimír Hanus (Čsl. akad. věd., Prague). *Chem. Listy* 50: 418-421 (1956). The polarographic reduction of phthalic acid on the dropping Hg-electrode in buffered solns. was accompanied by 3 waves, the reduction of corresponding monoesters by 2, and that of some diesters by 1 wave. The appearance and the height of the waves depended on the pH value of the soln. The interpretation of these waves was based on the assumption of various recombination possibilities, according to which the 1st of the waves was produced by the reduction of phthalic acid in the form of a cation, the 2nd one by the reduction of the undissoc. mol., whereas the last one was the wave of the univalent anion. The rate consts. of recombination and dissocn. were computed. F. Štráfelda

①

*[Handwritten signature]*

CZECHOSLOVAKIA / Physical Chemistry. Kinetics. Combustion.  
Explosions. Topochemistry. Catalysis.

B-9

Abs Jour : Ref Zhur - Khimiya, No 3, 1959, No. 7594

Author : IV. Janda, Jan  
V. Janda, J.; Hanus, V.; Obertik, J.

Inst : Not given

Title : Catalysts of Vinyl Chloride Synthesis. IV. Intermediate  
Reaction Product Formed During Synthesis in Presence of  
Water. V. Form of Inactivation of Mercury Catalyst

Orig Pub : Chem. zvesti, 1958, 12, No 1, 37-47; No 3, 155-162

Abstract : IV. Description of preparation and properties of hitherto  
unknown compounds that are formed during synthesis of vinyl  
chloride over mercury catalysts in the presence of  $H_2O$ .  
The formula of one of the compounds has been identified as  
di-(alpha, alpha'-chloromercury-beta, beta-dichlorethyl-  
dichloromercury-acetaldehyde)-mercury oxide. In contrast

Card 1/2

20

CZECHOSLOVAKIA / Physical Chemistry. Kinetics. Combustion.  
Explosions. Topochemistry. Catalysis.

"APPROVED FOR RELEASE: 09/19/2001

CIA-RDP86-00513R000617910003-4

Abs Jour : Ref Zhur - Khimiya, No 3, 1959, No. 7594

with current views it is shown in the paper that mercury  
chloride catalyzes the formation of vinyl chloride and  
some other reactions which take place during synthesis of  
vinyl chloride, even when the catalyst is not deposited  
on a substance of large specific surface.

V. The mercury catalyst used in the synthesis of vinyl  
chloride is poisoned mostly by excess  $C_2H_2$  and  $H_2O$  vapor.  
The only apparent reaction resulting in reduction of  
mercury salt to  $Hg_2Cl_2$ , or even to metallic mercury, and  
in formation of elemental carbon, is the reaction between  
acetylene,  $Hg_2Cl_2$  and water. Communication III see RZhKhim,  
1959, 3936. -- Part of the summary by the authors

Card 2/2

Z/009/60/000/011/001/001  
E112/E153

AUTHORS: Dolejšek, Z. Grubner, O. Hanuš, V. Kössler, I.  
Matyska, B. and Vodehnal, J.

TITLE: Analytical Control of Isoprene Rectification

PERIODICAL: Chemický průmysl, 1960, No. 11, pp. 571 - 575

TEXT: For the stereoscopic polymerization of isoprene, monomers of sufficiently high quality are essential. Purification of isoprene on a large scale is carried out by distillation processes. Technical isoprene contains various saturated and unsaturated hydrocarbons with 4, 5 or 6 carbons. Separation is accomplished by azeotropic distillation, adding acetaldehyde, propylene oxide, methyl formate, methanol, isopentane, isopropylamine, acetone, water or aqueous acetone as azeotropic agent. As the literature does not contain sufficient data about the boiling points of the different mixtures the authors have undertaken a study of the normal rectification of isoprene on efficient columns and have followed the concentrations of the different components in the various cuts. The effect of water and methyl alcohol as azeotropic agents was also considered.

Card 1/ 6

Z/009/60/000/011/001/001  
E112/E153

### Analytical Control of Isoprene Rectification

Two types of isoprene from different sources were investigated: 1) Soviet material, with 96% isoprene content, and 2) Czechoslovak material, prepared from isobutylene and formaldehyde, with 13% isoprene. The different distillation fractions were analysed by mass spectrography, infrared spectroscopy and gas chromatography, using thermoconductivity cells for detection. A chromatogram of sample B (Czechoslovak), e.g. first sample of condensate from still-head is shown (Fig.1), revealing 8 peaks and identified as follows: 1) isobutylene, not isolated in pure state but found in one fraction in an amount of 15% together with 85% 3-methylbutene-1; 2) and 3), peaks appertaining to butene-1 and butene-2 (confirmation of structure through mass spectrography); 4) 3-methylbutene-1 (this compound was isolated from one fraction in 99.5 purity and identified spectroscopically by comparison with data in the literature; 5) 2-methylbutene-1 (this compound was identified by comparison with literature data. It was obtained by fractional distillation in approximately 80% purity. It was also obtained by preparative

Card 2/6



Z/009/60/000/011/001/001  
E112/E153

#### Analytical Control of Isoprene Rectification

gas chromatography, and both samples proved identical);  
6) isoprene: standard prepared by fractional distillation in 99.98% purity and by preparative chromatographic method (ethyl cyclopentanecarboxylate as stationary phase); 7) 2-methylbutene-2 prepared by fractional distillation in 98% purity (identified by method used for 3-methylbutene-1; compound prepared for identification purpose also by preparative gas chromatography). Chromatogram of sample A (Soviet isoprene) revealed similar characteristics. A special peak (4b) was noticed, the identity of which was not yet determined. Results of practical distillation tests were as follows. Sample A was distilled over a low-efficiency column with reflux ratio 13:1. Pentene contents were reduced from 4 to 1.2%, and isoprene of 98.8% purity and in yields of 80% was collected. Using a more efficient column with reflux ratio 40:1 equilibrium was established after 2 hours and isoprene of 99.98% purity was obtained in poor yields. Attempts to improve yields by the addition of azeotropic agents (methanol, water) failed. Distillation of sample B was undertaken

Card 3/6

✓

Z/009/60/000/011/001/001  
E112/E153

# Analytical Control of Isoprene Rectification

over a column with reflux ratio 4:1. The concentration of isoprene in the middle fraction was doubled and the distillate contained only four components: 3-methylbutene-1; 2-methylbutene-1; isoprene; 2-methylbutene-2. A further fractionation over a column with reflux ratio 25:1 yielded further fraction, from which only those containing 2-methylbutene-1, isoprene and 2-methylbutene-2 were collected. Distillation of the three combined fractions over a column with reflux ratio 40:1 gave a two-component mixture in which the pentene concentration amounted to only 13%. By azeotropic distillation with acetone, conversion into high-grade isoprene could be achieved. It is claimed that yields were satisfactory. Acknowledgements are made to Doctor J. Pech, director, VÚSK Gottwaldov for useful advice and for supplying some of the raw materials. There are 6 figures, 4 tables and 16 references (including several patents to one reference): 11 English, 4 Czech and 1 Soviet.

ASSOCIATION: Ústav fyzikální chemie ČSAV, Praha (Institute for Physical Chemistry, ČSAV Prague)

Card 4/6

SUBMITTED: June 6, 1960

Z/009/60/009/011/001/001  
E112/E153

Analytical Control of Isoprene Rectification

Obr. 1  
*Chromatografická  
spektrum isoprenu B*

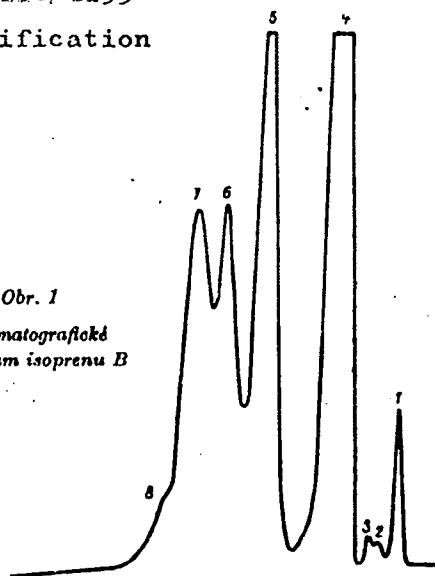


Fig.1

Card 5/6

Z/009/60/000/011/001/001  
E112/E153

Analytical Control of Isoprene Rectification

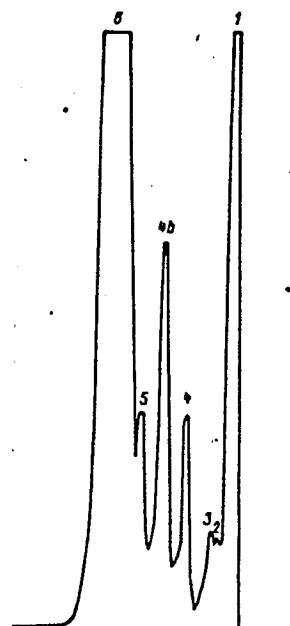


Fig.4

Card 6/6

Z/038/60/000/010/005/006  
A201/A026

AUTHORS: Hanůs, Vladimír; Dolejšek, Zdeněk

TITLE: Rearrangement of Some C<sub>7</sub>H<sub>8</sub> Isomers Induced by Electron-Impact Ionization

PERIODICAL: Jaderná energie, 1960, <sup>6</sup><sub>1</sub> No. 10, pp. 350 - 352

TEXT: S. Meyerson and P.N. Rylander (Ref. 1 through 4) have shown that ionization by electron impact in toluene results in the formation of tropylium ions. The authors carried further this study in an attempt to determine the influence of the structure of some other C<sub>7</sub>H<sub>8</sub> isomers on their behavior during ionization, and to learn the possibilities of some mass-spectrometric methods for the study of the structure and properties of ions in gaseous phase. This study has not been completed yet and the paper presents partial results obtained so far. The following C<sub>7</sub>H<sub>8</sub> isomers were studied: cycloheptatriene; toluene; spiro-(2.4)-heptadiene-(1.3); ethynyl cyclopentene-(1); ethynyl cyclopentene-(2); and bicyclo-(2.2.1)-heptadiene-(2.5). The following studied: the relative representation of fragmentation ions in dependence on the ionizing electron energy; the influence of the drawing-out voltage on the mass spectra; and the decomposition of accelerated ions,

Card 1/6

Z/038/60/000/010/005/006

A201/A026

Rearrangement of Some  $C_7H_8$  Isomers Induced by Electron-Impact Ionization

both spontaneous and induced by collisions with molecules of inert gases. Measurements were performed with a Nier-type mass spectrometer. It was found that the mass spectra of all six compounds, when obtained under similar conditions, were similar with a few exceptions in ethynyl cyclopentene-(1) and bicyclo(2.2.1)-heptadiene-(2.5). The spectra obtained are shown in Figure 1. Also changes of the electric field, by which ions are drawn out of the source, exerted practically the same influence on the mass spectra of all six compounds. It was further found that the definition properties of the instrument depend primarily on the kinetic energy of ions and not on their mass, and that with the exception of metastable ions the time during which ions remain in the source has no appreciable influence. In all compounds studied the same metastable ions were observed, corresponding to the spontaneous decomposition of ions during their time of flight from the accelerating field slit to the magnetic analyzer. These similarities indicate that the ionizing fragmentation in all six compounds proceeds basically in the same way. The main reaction, common to these isomers, is the formation of a molecular ion with a seven-membered cyclic structure, which decomposes to tropylium ions by splitting-off of a hydrogen atom. In ethynyl cyclopentene-(1), and even more in bicyclo-(2.2.1)-heptadiene-(2.5), competing side reactions of the decomposition of the excited

Card 2/6

Z/038/60/000/010/005/006  
A201/A026

Rearrangement of Some  $C_7H_8$  Isomers Induced by Electron-Impact Ionization

molecular ions assert themselves in addition to the above main reaction. In the former, the reaction competing with the formation of a fragmentation tropylium ion is the one by which  $C_5H_3^+$  is formed, which is then represented in a larger proportion than in cycloheptatriene. In bicyclo-(2.2.1)-heptadiene-(2.5), an increased rate of  $C_5H_6^+$ ,  $C_3H_4^+$  and  $C_3H_3^+$  ions can be observed. The large representation of  $C_5H_6^+$  ions obviously corresponds with the splitting-off of acetylene from the molecular ion. These ions do not decompose further to  $C_5H_3^+$ . The similarity of the spectra of the other four compounds indicates that in the rearranged parent ions, which for these compounds have the same structure, the excitation energy is distributed across the entire molecule in such a manner that equilibrium is reached before the fragmentation of parent ions takes place. The differences in the relative representation of molecular ions, tropylium ions, and the remaining fragmentation ions are related to the extent of the excitation energy in the rearranged parent ions, and are obviously due to the difference in the combination heat of the parent compounds. It is remarkable that from the total amount of ions, the portion of molecular ions depends on the combination heat to a far greater extent than the portion of ions formed by the tropylium fragmentation. The possibility cannot be

Card 3/6

Z/038/60/000/010/005/006  
A201/A026

Rearrangement of Some  $C_7H_8$  Isomers Induced by Electron-Impact Ionization

excluded that upon the formation of tropylium ions, or immediately thereafter, before they decompose any farther, an energy emission by radiation transitions takes place, which might reduce the original differences in the extent of the excitation energy of rearranged  $C_7H_8$  ions formed by ionization from different isomers. Should it become possible to prove that ions with the structure of parent moleculars practically are not represented among the molecular ions of these compounds, then the obtained data would largely contribute towards a more accurate understanding of processes preceding ionization fragmentation of more complicated molecules by electrons with energies in the range of several tens eV. There are 8 figures and 8 references: 7 English and 1 German.

ASSOCIATION: Ústav, fyzikální chemie ČSAV (Institute of Physical Chemistry, ČSAV)  
in Prague

Card 4/6

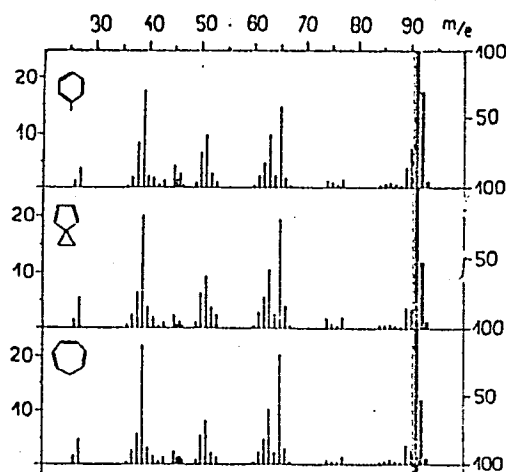


Z/038/60/000/010/005/006

A201/A026

Rearrangement of Some  $C_7H_8$  Isomers by Electron-Impact Ionization

Figure 1: Mass spectra of (A) toluene; (B) spiro-(2.4)-heptadiene-(1.3); (C) cycloheptatriene; (D) ethynyl cyclopentene-(2); (E) ethynyl cyclopentene-(1); (F) bicyclo(2.2.1)-heptadiene-(2.5)

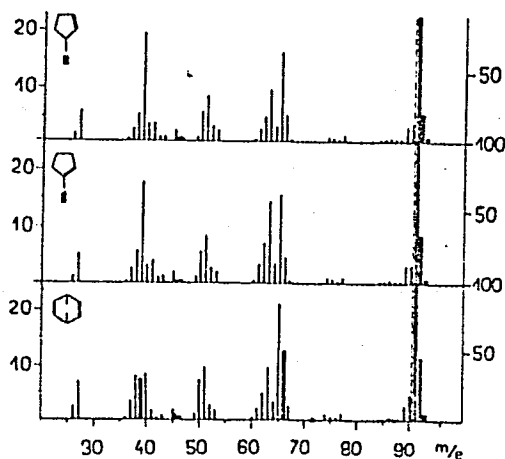


Card 5/6

Z/038/60/000/010/005/006  
A201/A025

Rearrangement of Some  $C_7H_8$  Isomers by Electron-Impact Ionization

(Figure 1 cont'd)



Card 6/6

AUTHORS: Koutecky, J., Hanuš, V.,  
Mayranovskiy, S. G. (Moscow)

S/076/60/034/03/025/038  
B005/B016

TITLE: Polarographic Catalytic Hydrogen Waves Caused by Organic Catalysts.  
I. Exact Solution of the Problem for the Case of a Bimolecular  
Regeneration of the Inactive Form of the Depolarizer From the  
Products of Electrode Reaction, and of a Monomolecular Conversion  
of the Depolarizer to Its Active Form

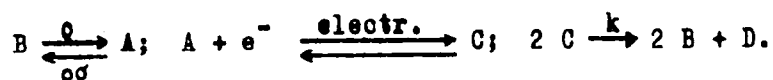
PERIODICAL: Zhurnal fizicheskoy khimii, 1960, Vol 34, Nr 3, pp 651-660 (USSR)

TEXT: In connection with investigations on catalytic hydrogen separation from  
buffer solutions on a dropping mercury electrode the authors aimed at explaining  
the complicated depolarization process for those cases in which the rates of the two  
chemical reactions taking place on the electrodes are very high (this means that  
equilibrium between diffusion and the chemical reactions is quickly established),  
and in which the concentration of the proton donors practically does not change  
with time. In this case, the concentrations of the components of the buffer system  
may be included into the two effective rate constants  $q$  and  $q_0$  of a monomolecular  
reversible reaction. The following schematic reaction equations are then obtained  
for the depolarization process:

Card 1/3

Polarographic Catalytic Hydrogen Waves Caused by Organic Catalysts. I. Exact Solution of the Problem for the Case of a Bimolecular Regeneration of the Inactive Form of the Depolarizer From the Products of Electrode Reaction, and of a Monomolecular Conversion of the Depolarizer to Its Active Form

S/076/60/034/03/025/038  
B005/B016



B - base (electrochemically inert form of the depolarizer); A - depolarizer (electrochemically active acid conjugated to B, which is in equilibrium with B in the solution); C - product of the electrode process; D - hydrogen. For the calculations an ideal mobility was assumed for the electrochemical equilibrium, so that the ratio  $\lambda = [A]_0/[C]_0$  depends only on the potential, but not on time. As a

further simplification the diffusion coefficients for the substances A, B, and C were assumed to be equal. It results therefrom that the algebraic sum of the diffusion currents flowing through random cross section is equal to zero. It was further assumed that the rates of all reactions mentioned above are very high already at a relatively small distance from the electrode,  $qt_1 \gg 1$ ;  $q\sigma t_1 \gg 1$ ;

$kat_1 \gg 1$ .  $t_1$  - dropping period of the mercury electrode;  $\alpha = [A] + [B] + [C]$ . Under these conditions a steady state forms after a very short time  $t_s$  of electrolysis  
Card 2/3